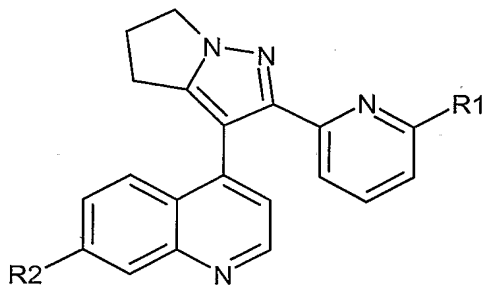


-102-

We claim:

1. A compound of the formula:



Formula I

wherein,

R1 represents hydrogen, halo, or (C1-C4)alkyl; and

R2 represents:

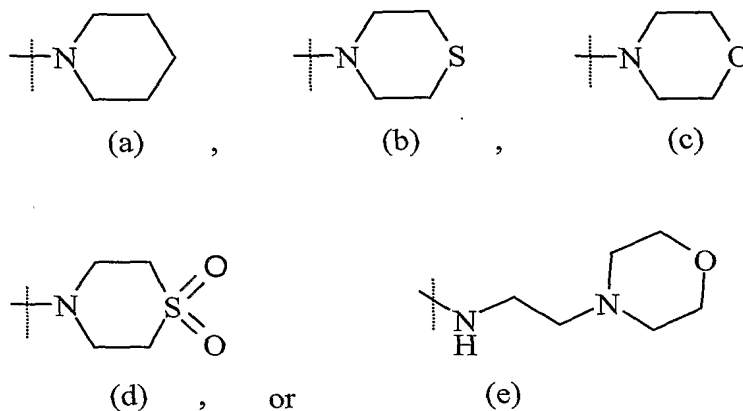
- (a) aryl;
- (b) aryl optionally substituted one to three times with a substituent independently selected from the group consisting of:
 - (i) halo,
 - (ii) amino,
 - (iii) nitro,
 - (iv) hydroxy,
 - (v) cyano,
 - (vi) (C₁-C₄)alkyl,
 - (vii) (C₁-C₄)alkoxy,
 - (viii) hydroxy(C₁-C₄)alkyl,
 - (ix) amino(C₁-C₄)alkyl,
 - (x) hydroxy(C₁-C₄)alkoxy,
 - (xi) halo(C₁-C₄)alkoxy,
 - (xii) (C₁-C₄)alkoxy(C₁-C₄)alkoxy,
 - (xiii) trifluoromethyl,
 - (xiv) difluoromethyl,
 - (xv) trifluoromethoxy,
 - (xvi) difluoromethoxy,
 - (xvii) (C₃-C₇)cylcoalkyl,
 - (xviii) COR³,

-103-

- (xix) (C₁-C₄)alkyl-COR⁴,
 (xx) amino(C₁-C₄)alkyl- COR⁴,
 (xxi) hydroxy(C₁-C₄)alkyl- COR⁴
 (xxii) (C₁-C₄)alkoxy-COR⁵,
 (xxiii) -C(NH₂)=N-OH
 (xxiv) NHSO₂R⁶,
 (xxv) SO₂R⁷,
 (xxvi) NHCOR⁸,
 (xxvii) SOR⁹,
 (xxviii) SR¹⁰,
 (xxix) CONHR¹¹,
 (xxx) O-(CH₂)_q-NR¹²R¹³, wherein q represents 1-4,
 (xxxi) tetrazole,
 (xxxii) methyltetrazole, and
 (xxxiii) CONCH-NR¹⁵R¹⁶
- (c) heterocycle;
 (d) heterocycle optionally substituted one to three times with a substituent independently selected from the group consisting of:
- (i) halo,
 (ii) amino,
 (iii) (C₁-C₄)alkyl,
 (iv) (C₁-C₄)alkoxy,
 (v) halophenyl(C₁-C₄)alkyl,
 (vi) (C₁-C₄)alkyl-(C₁-C₄)alkoxycarbonyl,
 (vii) CHO,
 (viii) COR³, and
 (ix) SO₂R⁷,
- (e) benzofused heterocycle;
 (f) benzofused heterocycle optionally substituted one or two times with a substituent independently selected from the group consisting of:
- (i) halo,
 (ii) amino,
 (iii) (C₁-C₄)alkyl,
 (iv) (C₁-C₄)alkoxy, and
 (v) (C₁-C₄)alkylcarbonyl,
 or (g) (C₃-C₇)cylcoalkyl;

-104-

R³ represents independently at each occurrence amino, hydroxy, (C₁-C₄)alkyl, (C₁-C₄)alkoxy, NH-(C₁-C₄)alkylamine, N,N-(C₁-C₄)dialkylamine, or a heterocycle selected from the group consisting of:



5

R⁴ and R⁵ represent independently at each occurrence amino, hydroxy, (C₁-C₄)alkyl, or (C₁-C₄)alkoxy;

R⁶ and R⁷ represent independently at each occurrence amino or (C₁-C₄)alkyl;

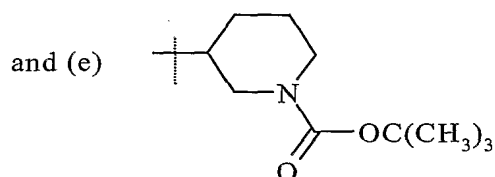
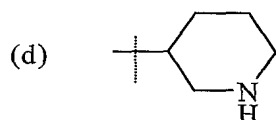
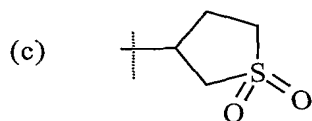
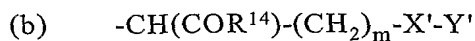
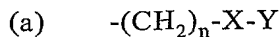
R⁸ represents independently at each occurrence amino, (C₁-C₄)alkyl, or (C₁-C₄)alkoxy;

10

R⁹ and R¹⁰ represent independently at each occurrence (C₁-C₄)alkyl;

R¹¹ represents independently at each occurrence (C₁-C₄)alkyl or a substituent selected from the group consisting of:

-105-



wherein,

n and m each independently represent 0-4;

5 X and X' represent independently at each occurrence $-\text{CO}-$, $-\text{CH}_2-$, $-\text{NH}-$, $-\text{S}-$, or $-\text{SO}_2-$; and

Y and Y' represent independently at each occurrence amino, hydroxy, (C_1-C_4) alkyl, (C_1-C_4) alkoxy, (C_1-C_4) alkoxycarbonyl, $\text{NH}-(\text{C}_1-\text{C}_4)$ alkylamine, or $\text{N,N}-(\text{C}_1-\text{C}_4)$ dialkylamine,

10 provided that where X or X' represents S, then Y or Y'' is not amino or hydroxy;

R^{12} and R^{13} represent independently at each occurrence hydrogen or (C_1-C_4) alkyl, or R^{12} and R^{13} together with the nitrogen atom to which they are attached form a piperidino, pyrrolidino, morpholino or a methylpiperazino group;

15 R^{14} represents independently at each occurrence hydroxy, amino, or (C_1-C_4) alkoxy; and

R^{15} and R^{16} each represent independently at each occurrence hydrogen or (C_1-C_4) alkyl,

or a pharmaceutically acceptable salt thereof.

20

-106-

2. A method of treating congestive heart failure comprising administering to a patient in need thereof an effective amount of the compound according to Claim 1.

3. A pharmaceutical composition comprising as an active ingredient a compound according to Claim 1 in combination with a pharmaceutically acceptable carrier, diluent or excipient.

4. The use of a compound according to Claim 1 for the manufacture of a medicament for the treatment of congestive heart failure.

10

15

20

25

30

35

40